Appendix Table 1. Primary Outcomes of Interest

| Quality of care outcomes | Outcome measures | | |
|---|---|--|--|
| Screening and other preventive care services completed or ordered | Screening for hypertension, hyperlipidemia, and diabetes in people not diagnosed with a CVD risk factor. Other preventive care services included providing aspirin (when appropriate), providing smoking cessation counseling, and nutrition and physical activity assessments included in USPSTF recommendations. ¹⁸⁻²² | | |
| | Clinical tests from evidence-based guidelines and protocols for management of hypertension, hyperlipidemia, or diabetes. | | |
| Clinical tests completed or ordered | Examples: hemoglobin A1C testing every 6 months for patients with diabetes ⁶ ; blood pressure and cholesterol testing for patients diagnosed with hypertension and hyperlipidemia, respectively. ^{7,8} | | |
| Treatments prescribed | Recommendations to initiate, intensify, or change existing medications for patients with hypertension, hyperlipidemia, or diabetes, based on clinical guidelines. | | |
| CVD risk factor outcomes | | | |
| Blood pressure outcomes | Proportion of patients with their BP controlled (usually defined as ≤140/90 mmHg and ≤130/80 mmHg for people with diabetes) ⁸ Change in mean SBP Change in mean DBP | | |
| Lipid outcomes | Proportion of patients achieving established targets (or better) for TC, LDL-cholesterol, HDL-cholesterol, and triglycerides ²³ Changes in mean TC, LDL-cholesterol, HDL-cholesterol, and triglycerides | | |
| Diabetes outcomes | Proportion of patients achieving A1C control (usually defined as $\leq 7\%$) ⁶ Changes in mean A1C level Changes in mean blood glucose levels | | |
| Other primary outcomes | | | |
| Morbidity, mortality, and patient- centered outcomes | Incidence of heart attacks and strokes CVD-related hospitalizations and ED visits Patient satisfaction with care Health-related quality of life | | |

BP, blood pressure; DBP, diastolic blood pressure; ED, emergency department; HDL, high-density lipoprotein; LDL, low-density lipoprotein; SBP, systolic blood pressure; TC, total cholesterol.

Appendix Table 2. Calculation of Individual Study Effect Estimates for Primary and Secondary Outcomes

| Effect estimate | Formula | | |
|--|---|--|--|
| Absolute percentage point change ^{a,b} | $(CDSS\ Prop_{post} - CDSS\ Prop_{pre}) - (UC\ Prop_{post} - UC\ Prop_{pre})$ | | |
| Difference-in-differences of the mean ^{a,b} | $(CDSS\ M_{post}-CDSS\ M_{pre})-(UC\ M_{post}-UC\ M_{pre})$ | | |

^aFor studies reporting multiple intervention arms, effect estimates were calculated for each arm and reported separately.

CDSS, clinical decision-support system; Prop, proportion of patients achieving desired outcome; Post, measurement from last available time point with ongoing CDSS; Pre, last measurement before intervention; UC, usual care; M, mean, average for patient group.

^bWhen studies reported multiple outcome measures (e.g., studies reporting multiple types of clinical tests), effect estimates for each measure were calculated and reported separately.

Appendix Table 3. Changes in Blood Pressure, Lipid, and Diabetes Outcomes Attributable to Clinical Decision-Support Systems

| Outcome type | Number of studies | Median effect estimate (IQI) |
|--|--|---------------------------------|
| Blood pressure outcomes | | |
| Improvement in proportion of patients | 8 ^{25,33,39,41,52,63,65,82} | +2.0 pct pts |
| with BP at goal ^a | | (-5.0, 10.5) |
| Reduction in SBP | $14^{25,26,37-39,41,45,50,51,58,63,65,83,85}$ | –0.89 mmHg |
| | | (-1.93, 1.0) |
| Reduction in DBP | 11 ^{26,38,39,41,45,50,51,65,71,83,85} | –0.30 mmHg |
| | | (-1.13, 1.0) |
| Lipid outcomes | | |
| Improvement in proportion of patients | 9 ^{26,33,41,45,65,81,82,85,92} | +1.0 pct pts |
| with lipid at goal ^b | | (-1.25, 4.55) |
| Reduction in total cholesterol | 5 ^{26,27,37,38,84} | 0 mg/dL |
| | | (-7.35, 4.4) |
| Reduction in LDL cholesterol | $10^{26,27,29,41,45,58,65,83-85}$ | −0.5 mg/dL |
| | | (-2.4, 0.2) |
| Improvement in HDL cholesterol | 3 ^{26,27,84} | −0.27 mg/dL |
| | | (NA) |
| Reduction in triglycerides | $2^{27,84}$ | −21.4 mg/dL |
| | | (NA) |
| Diabetes outcome | | |
| Improvement in the proportion of | 8 ^{26,33,41,45,65,82,85,92} | -1.3 pct pts |
| patients with A1C at goal ^c | | (-2.15, 4.23) |
| Reduction in A1C level | 11 ^{26,29,38,41,45,52,58,65,83-85} | -0.12% |
| | | (-0.28, 0) |

^aAbsolute percentage point increase in proportion of patients achieving goal BP.

BP, blood pressure; DBP, diastolic blood pressure; HDL, high-density lipoprotein; IQI, interquartile interval; LDL, low-density lipoprotein; NA, not applicable; pct pts, percentage points; SBP, systolic blood pressure.

^bAbsolute percentage point increase in proportion of patients achieving goal lipid levels.

^cAbsolute percentage point increase in proportion of patients achieving goal A1c levels.

Appendix Table 4. Changes in Quality of Care Outcomes from Studies Examining Clinical Decision-Support Systems Combined with Other Interventions

| Additional intervention delivered | Quality of care outcome | Study author (year) | Quality of care findings ^a |
|-----------------------------------|---|----------------------------------|---|
| Team-based care | Screening and other preventive care services completed or ordered | Holbrook (2011) ⁸³ | • Change in total process composite score (95% CI): +4.7 (3.63 to 5.71) |
| | | Dorr (2005) ²⁹ | HbA1c testing completed: OR (95% CI): 1.5 (1.3 to 1.7) LDL testing completed: OR (95% CI): 1.3 (1.0 to 1.6) |
| | Clinical tests completed or ordered | O'Connor (2005) ⁵² | Prop of patients with ≥2 HbA1c tests: +33.0 pct pts (p=0.002) Prop of patients with ≥1 LDL test: +16.0 pct pts (p=0.12) Prop of patients with ≥2 HbA1c tests and with ≥1 LDL test: +25.0 pct pts (p=0.03) |
| | | Hicks (2008) ³⁹ | • Prop of patients with recommended drug class prescribed: +2.0 pct pts (<i>p</i> <0.001) |
| | Treatments prescribed | Murray (2004) ⁵¹ | Prop of patients prescribed antihypertensive medications (95% CI) Arm 1: Pharmacist + CDSS vs. UC –2.0 pct pts (–1.90 to 7.90) Arm 2: Pharmacist + MD + CDSS vs. UC: +2.0 pct pts (–7.80 to 11.80) |

| Screening and other Patient reminders preventive care services completed or ordered | Holbrook (2009) ⁴¹ | Change in BP process composite score: +0.61 (p<0.001) Change in BMI process composite score: +0.71 (p<0.001) Change in exercise process composite score: +0.91 (p<0.001) Change in diet process composite score: +0.88 (p<0.001) Change in aspirin process composite score: +0.05 (p=0.02) Change in smoking process composite score: +0.03 (p=0.09) | |
|---|----------------------------------|---|---|
| | | Ornstein (1991) ⁵³ | Prop of patients receiving cholesterol screening Arm 1: CDSS vs UC: +9.10 pct pts (p<0.001) Arm 2: CDSS + patient reminder vs UC: +18.6 pct pts (p<0.001) |

^aFindings are individual effect estimates (absolute percentage point difference or difference-in-differences of the mean or odds ratios) on the effectiveness of a CDSS intervention compared with usual care.

BP, blood pressure; CDSS, clinical decision-support system; pct pts, percentage points; Prop, proportion; UC, usual care

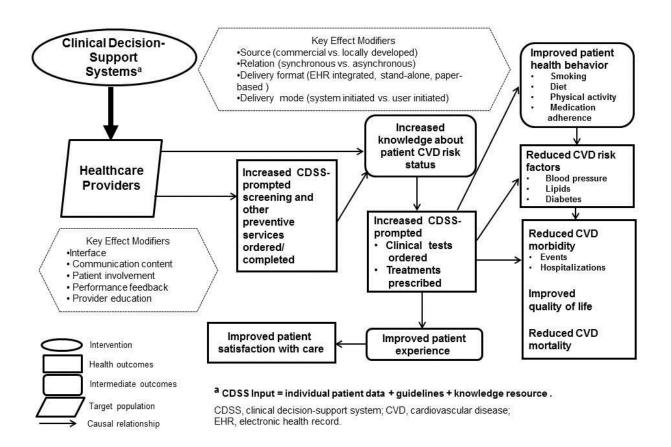
Appendix Table 5. CDSS Contextual Factors and Features for Health Process Outcomes

| Contextual factor/feature | Number of studies reporting (% of included studies) N=45 | Number of studies reporting favorable outcome measures for screening and other preventive services (% of total studies reporting outcome) n=17 | Number of studies reporting favorable outcome measures for clinical tests ordered (% of total studies reporting outcome) n=7 | Number of studies reporting favorable outcome measures for treatments prescribed (% of total studies) n=11 |
|---|--|--|--|--|
| Integration with charting or order entry system to support workflow integration | 33 (73.3) | 9 (52.9) | 3 (42.9) | 5 (45.5) |
| Automatic provision of decision support as part of clinician workflow | 38 (84.4) | 10 (58.8) | 4 (57.1) | 6 (54.5) |
| No need for additional clinician data entry | 24 (53.3) | 6 (35.3) | 2 (28.5) | 5 (45.5) |
| Request documentation of the reason for not following CDSS | 4 (8.9) | 2 (11.8) | 1 (14.3) | 0 (0) |
| Provision of decision support at time and location of decision making | 35 (77.8) | 9 (52.9) | 3 (42.9) | 6 (54.5) |
| Recommendations executed by noting agreement | 3 (6.7) | 2 (11.8) | 0 (0) | 1 (9.1) |
| Provision of a recommendation not just an assessment | 41 (91.1) | 10 (58.8) | 5 (71.4) | 7 (63.6) |
| Promotion of action rather than inaction | 15 (33.3) | 4 (23.5) | 1 (14.3) | 3 (27.2) |
| Justification of decision support via provision of reasoning | 8 (17.8) | 1 (5.9) | 0 (0) | 1 (9.1) |
| Justification of decision support via provision of research evidence | 11 (24.4) | 1 (5.9) | 1 (14.3) | 1 (9.1) |
| Local user involvement in development process | 18 (40) | 2 (11.8) | 2 (28.6) | 1 (9.1) |

Appendix
Clinical Decision Support Systems and Prevention: A Community Guide Cardiovascular Disease Systematic Review
Njie et al.

| Provision of decision support results to patients as well as providers | 8 (17.8) | 2 (11.8) | 1 (14.3) | 1 (9.1) |
|--|-----------|----------|----------|----------|
| CDSS accompanied by periodic performance feedback | 11 (22.9) | 2 (11.8) | 1 (14.3) | 2 (18.1) |
| CDSS accompanied by conventional education | 12 (25) | 4 (23.5) | 0 (0) | 3 (27.2) |

Appendix Figure 1. Analytic framework: clinical decision-support systems for cardiovascular disease prevention.



Appendix Figure 2. Search process.

